IN THE CLAIMS:

Cancel claims 1-4, 6-8, and 27 without prejudice or disclaimer.

Please amend claims 5 and 9 and add new claims 30-52 as shown in the below LISTING OF CLAIMS.

Claims 1-4 (canceled)

Claim 5 (currently amended): <u>An isolated The polynucleotide according to claim 2,</u> comprising the nucleic acid sequence as shown in of SEQ ID No. NO: 1.

Claims 6-8 (canceled)

Claim 9 (currently amended): An isolated A polynucleotide sequence according to claim 1, wherein the polynucleotide codes for which encodes a polypeptide that comprises the amino acid sequence shown in of SEQ ID NO: 2.

Claim 10 (withdrawn): A coryneform bacteria in which the deaD gene is attenuated.

Claim 11 (withdrawn): The coryneform bacteria according to claim 10, wherein the deaD gene is eliminated.

Claim 12 (original): An Escherichia coli strain Top10/pXK99EdeaD deposited as DSM 14464.

Claims 13 (withdrawn): A method for the fermentative preparation of L-amino acids in coryneform bacteria, comprising:

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a) fermenting, in a medium, the coryneform bacteria which produce the desired L-amino acid and in which at least the deaD gene or nucleotide sequences which code for it are attenuated.

Claims 14 (withdrawn): The method according to claim 13, further comprising:

b) concentrating the L-amino acid in the medium or in the cells of the bacteria.

Claims 15 (withdrawn): The method according to claim 14, further comprising:

c) isolating the L-amino acid.

Claims 16 (withdrawn): The method according to claim 13, wherein the L amino acids are lysine.

Claims 17 (withdrawn): The method according to claim 13, wherein deaD gene or nucleotide sequences coding for this gene are overexpressed.

Claims 18 (withdrawn): The method according to claim 13, wherein additional genes of the biosynthesis pathway of the desired L-amino acid are enhanced in the bacteria.

Claims 19 (withdrawn): The method according to claim 13, wherein bacteria in which the metabolic pathways which reduce the formation of the desired L-amino acid are at least partly eliminated are employed.

Claims 20 (withdrawn): The method according to claim 13, wherein the expression of the polynucleotide(s) which code(s) for the deaD gene is attenuated.

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Claims 21 (withdrawn): The method according to claim 20, wherein the expression of the polynucleotide(s) which code(s) for the deaD gene is eliminated.

Claims 22 (withdrawn): The method according to claim 13, wherein the catalytic properties of the polypeptide for which the polynucleotide deaD codes are reduced.

Claims 23 (withdrawn): The method according to claim 13, wherein the bacteria being fermented comprise, at the same time, one or more genes which are enhanced or overexpressed; wherein the one or more genes is/are selected from the group consisting of:

the dapA gene which codes for dihydrodipicolinate synthase,

the gap gene which codes for glyceraldehyde 3-phosphate dehydrogenase,

the tpi gene which codes for triose phosphate isomerase,

the pgk gene which codes for 3-phosphoglycerate kinase,

the zwf gene which codes for glucose 6-phosphate dehydrogenase,

the pyc gene which codes for pyruvate carboxylase,

the mqo gene which codes for malate-quinone oxidoreductase,

the lysC gene which codes for a feed-back resistant aspartate kinase,

the lysE gene which codes for lysine export,

the hom gene which codes for homoserine dehydrogenase

the ilvA gene which codes for threonine dehydratase or the ilvA(Fbr) allele which codes for a feed back resistant threonine dehydratase,

the ilvBN gene which codes for acetohydroxy-acid synthase, the ilvD gene which codes for dihydroxy-acid dehydratase, and the zwa1 gene which codes for the Zwa1 protein.

Claims 24 (withdrawn): The method according to claim 13, wherein the bacteria being fermented comprise, at the same time, one or more genes which are attenuated; wherein the genes are selected from the group consisting of:

the pck gene which codes for phosphoenol pyruvate carboxykinase, the pgi gene which codes for glucose 6-phosphate isomerase, the poxB gene which codes for pyruvate oxidase, and the zwa2 gene which codes for the Zwa2 protein.

Claims 25 (withdrawn): The method according to claim 13, wherein microorganisms of the species Corynebacterium glutamicum are employed.

Claims 26 (withdrawn): The method according to claim 25, wherein the Corynebacterium glutamicum strain DSM5715::pXK99EdeaD is employed.

Claim 27 (canceled)

Claim 28 (withdrawn): A method for discovering RNA, cDNA and DNA in order to isolate nucleic acids or polynucleotides or genes which code for DNA/RNA helicase or have a high similarity with the sequence of the deaD gene, comprising contacting the RNA, cDNA, or DNA with hybridization probes comprising polynucleotide sequences according to claim 1.

Claim 29 (withdrawn): The method according to claim 28, wherein arrays, micro arrays or DNA chips are employed.

Claim 30 (new): An isolated polynucleotide that is at least 90% identical to SEQ ID NO: 1 and encodes a polypeptide that has the enzymatic activity of a DNA/RNA helicase.

Claim 31 (new): The isolated polynucleotide of claim 30, wherein said polynucleotide is at least 95% identical to SEQ ID NO: 1.

Claim 32 (new): The isolated polynucleotide of claim 30, wherein said polynucleotide is at least 99% identical to SEQ ID NO: 1.

Claim 33 (new): An isolated polynucleotide which encodes a polypeptide comprising the amino acid sequence of SEQ ID NO: 2, wherein said polypeptide has the enzymatic activity of a DNA/RNA helicase.

Claim 34 (new): An isolated polynucleotide comprising nucleotides 259 to 2130 of SEQ ID NO: 1.

Claim 35 (new): An isolated polynucleotide consisting of SEQ ID NO: 1 or a fragment of SEQ ID NO: 1 that encodes a polypeptide having the enzymatic activity of a DNA/RNA helicase.

Claim 36 (new): An isolated polynucleotide that hybridizes to the complete complement of SEQ ID NO: 1 under stringent conditions comprising a final wash at 68°C, wherein said isolated polynucleotide encodes a polypeptide that has the enzymatic activity of a DNA/RNA helicase.

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Claim 37 (new): An isolated polynucleotide comprising the nucleotide sequence of the complete complement of SEQ ID NO: 1.

Claim 38 (new): A vector comprising the isolated polynucleotide of any of the claims 30 or 33 to 37.

Claim 39 (new): An isolated polynucleotide consisting of at least 18 consecutive nucleotides of SEQ ID NO: 1 or the complete complement of SEQ ID NO: 1.

Claim 40 (new): The isolated polynucleotide of claim 39, wherein said polynucleotide consists of at least 20 consecutive nucleotides.

Claim 41 (new): The isolated polynucleotide of claim 30, wherein said polynucleotide consists of at least 18 consecutive nucleotides.

Claim 42 (new): The isolated polynucleotide of claim 41, wherein said polynucleotide consists of at least 20 consecutive nucleotides.

Claim 43 (new): A vector comprising the isolated polynucleotide of claim 39.

Claim 44 (new): The vector of claim 43, wherein said vector is pXK99EdeaD deposited in Escherichia coli Top/pXK99EdeaD under DSM 14464.

Claim 45 (new): A primer for the synthesis of a polynucleotide in a polymerase chain reaction comprising a DNA fragment consisting of at least 18 consecutive nucleotides of SEQ ID NO: 1 or the complete complement of SEQ ID NO: 1, wherein said

polynucleotide encodes a polypeptide that has the enzymatic activity of a DNA/RNA helicase.

Claim 46 (new): The primer of claim 45, wherein said DNA fragment consists of at least 20 consecutive nucleotides.

Claim 47 (new): A probe for the detection or isolation of a polynucleotide in a hybridization reaction comprising a DNA fragment consisting of at least 18 consecutive nucleotides selected from SEQ ID NO: 1 or the complete complement of SEQ ID NO: 1, wherein said polynucleotide encodes a polypeptide that has the enzymatic activity of a DNA/RNA helicase.

Claim 48 (new): The probe of claim 47, wherein said DNA fragment consists of at least 20 consecutive nucleotides.

Claim 49 (new): A recombinant host cell comprising the isolated polynucleotide of claims 30 or 33 to 37.

Claim 50 (new): The host cell of claim 49, wherein said host cell is of the species Escherichia coli.

Claim 51 (new): A recombinant host cell of the genus Corynebacterium or of the species Escherichia coli comprising the vector of claim 43.

Claim 52 (new): The host cell of claim 51, wherein said host cell is of the species Corynebacterium glutamicum.